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(54) Title: METHODS OF IDENTIFYING OPTIMAL VARIANTS OF PEPTIDE EPITOPES

Amino Acid Sequence	Binding IC <sub>50</sub> (nM)	# Isolates	Predicted Cross-reactivity		Immunogenicity (SU)
			MTNPPPIPV	MTSNPPPIPV	
MTSNPPPIPV	52.8	60	-	+	10
MTNPPPIPV	128.4	33	+	+	100
MTSNPPVPV	21.8	26	-	+	1000
MTGNPPPIPV	125.1	15	-	+	10000
MTGNPPVPV	2021	9	-	+	10
MTNPPVPV	85.6	6	+	+	100
MTANPPVPV	20.0	3	-	+	1000
MTHNPPPIPV	167.0	2	+	-	100
MTANPPPIPV	2.3	1	-	+	1000
MTSDPPPIPV	107.4	1	-	+	100
MTGNPSPVPV	15.8	1	-	+	100
MTGNPAIPV	1200	1	-	+	100
MTSNPAIPV	1465	1	-	+	100
MTRNPPVPV	9171	1	-	-	100

(57) Abstract: The present invention is directed to methods for selecting a variant of a peptide epitope which induces a CTL response against another variant(s) of the peptide epitope, by determining whether the variant comprises only conserved residues, as defined herein, at non-anchor positions in comparison to the other variant(s). The present invention is also directed to variants identified by the methods above; peptides comprising such variants; nucleic acids encoding such variants and peptides; cells comprising such variants, and/or peptides, and/or nucleic acids; compositions comprising such variants, and/or peptides, and/or nucleic acids, and/or cells; as well as therapeutic and diagnostic methods for using such variants, peptides, nucleic acids, cells, and compositions.



TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

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